

10/562181

IAP9 Rec'd PCT/PPO 22 DEC 2003

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(81) *Designierade stater* AP: all, AL, AM, AT, AU, AZ, BA, BB, BG, BR,
Designated states BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE,
DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE,
GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG,
KP, KR, KZ, LC, LI, LK, LR, LS, LT, LU, LV,
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SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA,
UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

(21) *Patentansökningsnummer* PCT SE04/00741
Patent application number

(86) *Ingivningsdatum* 2004-05-13
Date of filing

**CERTIFIED COPY OF
PRIORITY DOCUMENT**

Stockholm, 2005-10-03

För Patent- och registreringsverket
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Gunilla Larsson

Avgift
Fee 170:-

PCT REQUEST

Original (for SUBMISSION)

0	For receiving Office use only International Application No.	PCT / SE 2004 / 000741
0-2	International Filing Date	13 -05- 2004
0-3	Name of receiving Office and "PCT International Application"	The Swedish Patent Office PCT International Application
0-4	Form PCT/RO/101 PCT Request Prepared Using	PCT-SAFE [EASY mode] Version 3.50 (Build 0002.162)
0-5	Petition The undersigned requests that the present international application be processed according to the Patent Cooperation Treaty	
0-6	Receiving Office (specified by the applicant)	Swedish Patent Office (RO/SE)
0-7	Applicant's or agent's file reference	P 04-144
I	Title of Invention	DETECTION OF DIASTOLIC HEART FAILURE
II	Applicant II-1 This person is II-2 Applicant for II-4 Name II-5 Address	applicant only all designated States except US ST JUDE MEDICAL AB 175 84 JÄRFÄLLA Sweden
II-6	State of nationality	SE
II-7	State of residence	SE
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IAP9 Recd PCTPTO 22 DEC 2003

DETECTION OF DIASTOLIC HEART FAILURE

Technical Field

The present invention relates to an implantable medical apparatus for detecting diastolic heart failure, DHF, comprising a DHF determining device for determining at least one blood pressure parameter for detecting a DHF state of the heart of a patient. The invention also relates to a pacemaker provided with such an apparatus, and a method of detecting DHF, comprising the step of determining at least one blood pressure parameter for detecting a DHF state of the heart of a patient.

Background

There is a growing recognition that congestive heart failure caused by a predominant abnormality in the diastolic function, i.e. diastolic heart failure, DHF, is both common and causes significant morbidity and mortality. Therefore early detection of DHF is important. Patients do not, however, seem to have symptoms at an early stage. In addition it has been hard to separate diastolic and systolic heart failure and they may also exist simultaneously.

It has been discovered that among the few parameters, separating diastolic heart failure from systolic heart failure, are certain blood pressure parameters obtained during work of the patient. Thus US 6 438 408 describes an implantable medical device for monitoring congestive heart failure, CHF. A plurality of heart failure parameters indicative of the state of the heart failure are measured employing EGMs, blood pressures including absolute pressures, developed pressures (= systolic pressures – diastolic pressures) and the time derivative dP/dt, as well as heart chamber volumes. One of these parameters is the relaxation or contraction time constant τ of the heart chamber. τ is calculated from a continuous pressure signal and is the drop in ventricular pressure at the end of systole and in the first part of diastole. The τ parameter is thus a general parameter reflecting the relaxation process.

Patients suffering from DHF have both elevated ventricular filling pressure and elevated pulmonary venous pressure. To maintain cardiac output under load higher filling pressures are required to achieve adequate left ventricular filling, see H. Weinberger, Diagnosis and Treatment of Diastolic Heart Failure, Hospital

Practice, <http://www.hosppract.com/issues/1999/03/weinb.htm>. However, for DHF patients increased filling pressure does not result in increased left ventricular diastolic volumes. This is illustrated in figure 1 which shows results of studies on patients with isolated diastolic dysfunction, see D. W. Kitzman et al, "Exercise

5 Intolerance in Patients with Heart Failure and Preserved Left Ventricular Systolic Function: Failure of the Frank-Starling Mechanism", J. Am. Coll. Cardiol., 17:1065-1072, 1991. Figure 1a shows the pulmonary capillary wedge pressure, PCWP, as a function of workload for patients with isolated diastolic dysfunction and for control persons not suffering for this dysfunction, and figure 1b shows
10 corresponding curves for the left ventricular end-diastolic volume index, LVEDVI as a function of the workload. From the figures it appears that for patients suffering from the mentioned diastolic dysfunction, PCWP was elevated at rest and further elevated during exercise. This indicates that the ventricular filling pressure was increased under both conditions, whereas LVEDVI did not increase with exercise.

15 The present invention is based on the above-mentioned knowledge. The purpose of the present invention is consequently to utilize this teaching to propose a technique for detecting DHF based on measurement of left atrial or pulmonary vein pressure.

20 Disclosure of the Invention

This purpose is obtained by an apparatus, a pacemaker and a method of the kind mentioned in the introductory portion of the description and having the characterizing features of claims 1, 2, 12 , 16 and 17 respectively.

Thus with the present invention the reduced peak and submaximal exercise performance of DHF patients is utilized for detecting DHF. With a sensor placed in left atrium or very close to its endocardium the left atrial pressure is measured during the diastasis phase just before atrial contraction. Alternatively a sensor in the pulmonary vein can be used, provided that pressure data for subsequent analysis are collected only when the pulmonary valve is closed. With
25 the technique according to the invention it is possible to detect DHF at an early stage, often before the patient seems to have any symptoms.

In the present invention the workload situation of the patient must be identified, and therefore, according to an advantageous embodiment of the

apparatus according to the invention, an activity sensor is provided for determining the workload of the patient.

Since the absolute pressure is used, measured pressure values must be corrected with e.g. the level of the mean pressure in the right atrium or in vena cava close to the right atrium. This correction is needed because any column of blood above the sensor location, which thus is posture depending, will add to the absolute pressure level. Such a correction will also compensate for a varying ambient pressure. According to an advantageous embodiment of the apparatus according to the invention pressure compensating means are therefore provided for correcting the measured absolute value of left atrial pressure with the mean pressure in right atrium or in vena cava close to the right atrium.

According to another advantageous embodiment of the apparatus according to the invention the DHF determining device comprises an impedance measuring means for measuring the impedance between a left ventricular electrode and a second electrode intended to be positioned in such a place that most of variations in the measured impedance are due to varying left ventricular volume, and in that ΔP values and corresponding values of the difference, ΔV , in the end-diastolic ventricular volume obtained from said impedances measured for said predetermined workload situation and for the patient in rest are supplied to a quotient forming means to form the quotient $\Delta P/\Delta V$, and in that said comparison means is adapted to compare said quotient $\Delta P/\Delta V$ with a predetermined quotient reference value for DHF detection. By adding information about left ventricular end-diastolic pressure the DHF detection can be improved. In a healthy person the end-diastolic volume increases with exercise but shows only a small increase in left atrial pressure. In a person suffering from DHF, who has a more stiff heart muscle, the left ventricular end-diastolic volume changes very little but there is a large increase in left atrial pressure. The quotient $\Delta P/\Delta V$ is consequently a useful quantity for improving the detection of DHF. The difference ΔV is determined by impedance measurements.

According to still other advantageous embodiments of the apparatus according to the invention a first averaging means is provided to form an average value of the absolute pressures measured during a plurality of cardiac cycles with said predetermined workload situation and an average value of the absolute

pressures measured during a plurality of cardiac cycles with the patient in rest, and a second averaging means is provided to form an average value of said impedance measured during a plurality of cardiac cycles with said predetermined workload situation and an average value of said impedance measured during a plurality of cardiac cycles with the patient in rest. In this way the quality of the pressure measurements as well as the impedance measurements are improved.

According to other advantageous embodiments of the apparatus according to the invention a wireless communication means is connected to said comparison means for automatically sending the results of the comparisons with said reference values to external receiver means, or a storing means is provided for storing the results of the comparisons with said reference values. Thus if the measured absolute pressure, and possibly also the quotient $\Delta P/\Delta V$, have risen above the associated reference values in a predetermined way this condition is automatically transmitted to a physician or stored for transmission in connection with a follow-up.

The invention also relates to a pacemaker provided with the apparatus for detecting DHF and control means for optimising pacing therapy depending on the result of the comparisons with the predetermined reference values. The pressure measuring means of the apparatus according to the invention then preferably comprises a pressure sensor connected to the pacemaker, since it can monitor the pulse pressure of its carrier for long periods. This is an advantage since evolvement of DHF is a slow process.

An advantageous embodiment of the pacemaker according to the invention comprises a rate responsive sensor for use as activity sensor for determining the workload situation of the patient. Even the pressure sensor of the pacemaker can be used as activity sensor.

According to an advantageous embodiment of the method according to the invention in a calibration procedure instantaneous values of the left ventricular volume are measured by ultrasound technique and the relation is established between these measured ventricular volume values and simultaneously measured impedance values. Since the measured impedance depends on the location of the electrodes used, it will vary from patient to patient and therefore such a calibration is required.

As mentioned above the measured difference between left atrial pressure measured in workload and rest situations is compared with a predetermined pressure difference reference value. The reference value for detection of DHF is preferably obtained from measurements on the patient at an early stage of the 5 implantation period of the apparatus or pacemaker. The patient is supposed not to suffer from DHF at the time of implantation. Therefore, according to an advantageous embodiment of the method according to the invention, the pressures are measured for different workloads of the patient and for the patient in rest at an early time, when the patient is not suffering from DHF, for determining 10 the pressure difference reference value. Corresponding impedance measurements are preferably also made on the patient at this early stage for determination of the quotient reference value.

Brief Description of the Drawings

15 To explain the invention in greater detail an embodiment of the invention will now be described with reference to the drawings on which figures 1a and 1b are diagrams showing PCWP and LVEDVI respectively as functions of workload for patients having a diastolic dysfunction and control persons not suffering from this dysfunction, figure 2 is a plot of PCWP versus left ventricular end-diastolic 20 volume, figure 3 shows a block diagram of an embodiment of a pacemaker according to the invention, figure 4 is a flow chart illustrating a procedure for determination of the quotient $\Delta P/\Delta V$ in an embodiment of the apparatus according to the invention, and figure 5 is a flow chart illustrating an averaging process of the procedure illustrated in figure 4.

25

Description of Preferred Embodiments

Figures 1a and 1b show results of studies on patients having a diastolic dysfunction as explained above. From these figures it appears that a heart suffering from DHF has another PCWP versus workload behaviour than a normal 30 heart, "Controls" in the figure. The present invention is based on this knowledge since the PCWP is approximately equal to the left atrial pressure LAP.

The detection of DHF by measurements of the absolute left atrial pressure LAP or the pressure in the pulmonary vein can be improved by adding information about left ventricular end-diastolic volume. In a healthy person the end-diastolic

volume increases with workload or exercise, whereas the left atrial pressure shows only a small increase. In a person suffering from DHF the left ventricular end-diastolic volume changes vary little but there is a large increase in the left atrial pressure. This is illustrated by figure 2, which is a diagram from D. W.

5 Kitzman et al, "Exercise Intolerance in Patients with Heart Failure and Preserved Left Ventricular Systolic Function: Failure of the Frank-Starling Mechanism", J. Am. Coll. Cardiol., 17:1065-1072, 1991, plotting PCWP versus left ventricular end-diastolic volume. The open boxes in the figure indicate the direction of change from rest to exercise or workload for patients with DHF, and the solid boxes
10 illustrate the corresponding direction of change for normal control persons. The shift upwards in the diagram for sick patients should be noted.

The left ventricular end-diastolic volume is preferably determined by measurements of the electric impedance between e.g. a left ventricular electrode, normally placed in a coronary vein, and an electrode in right ventricle, right atrium
15 or any other place in or on the heart, such that most of the variations in the measured impedance derive from variations in the left ventricular volume, as will be described below.

The detection and measurement of degree of DHF can be improved by studying the quotient $\Delta P/\Delta V$. ΔP denotes the difference between left atrial
20 pressure measured in the workload and rest situations and ΔV the difference in the end-diastolic ventricular volume, obtained by the impedances measurements, for the predetermined workload situation and for the patient in rest, as explained above.

The quantity ΔV , and consequently the quotient $\Delta P/\Delta V$, is depending on
25 the positions of the electrodes, used for the impedance measurements. These quantities will therefore vary from patient to patient. Thus some kind of calibration is needed.

The left ventricular volume varies during a cardiac cycle. The instantaneous volume can be measured by e.g. ultrasound technique. By
30 simultaneously measuring corresponding impedance values a relation can be established between measured impedance values and left ventricular volume.

The measured pressure difference ΔP can be directly obtained since the sensitivity of the pressure transducer is set by the manufacturer.

The reference value for the quotient $\Delta P/\Delta V$ for detection of DHF can be set from common knowledge about physiological properties during the progression of DHF. In the same way the severity of the progression of DHF can be followed.

Figure 3 is a block diagram of an embodiment of a pacemaker according 5 to the invention, including a pressure sensor 2, sensitive to the pressure in left atrium, LAP, of a patient's heart 4. The atmospheric pressure and the pressure from blood volumes above the sensor also contribute to LAP. To compensate for this a sensor 6 is also provided for measuring the pressure in the right atrium or in vena cava. This compensation pressure is denoted by Pref.

10 A measurement value correlated to left ventricular volume changes ΔV is obtained by measuring the impedance Z between an electrode 8 in or attached to the left ventricle and an electrode 10 in the right ventricle. The parameters in question are measured in the diastasis phase just before the atrial contribution to the left ventricular filling. To do this a timer can be used, starting at the preceding 15 contraction. Alternatively P-waves can be sensed by an electrode 12 implanted in the right atrium of the heart 4 for starting the measurement. Parameter values measured during a period preceding the atrial contraction are stored in a ring buffer. This procedure will be described below with reference to figure 4.

20 The pacemaker in figure 3 comprises ordinary circuits 14 for IEGM sensing and stimulation. It also comprises circuitry 16 for measuring the impedance Z between implanted electrodes 8, 10 as mentioned above. To determine volume changes of heart chambers by impedance measurements between electrodes implanted in the heart is per se previously known from e.g. EP 1 384 492. Such impedance measuring circuitry is also known from respiration 25 activity measuring pacemakers.

The pacemaker illustrated in figure 3 also contains a unit 20 for measuring activity or workload of the patient by means of an activity or workload sensor 22. An ordinary microprocessor with supporting circuits 18 are provided for calculation and signal processing explained in the following with reference to figures 4 and 5. 30 Sensed Pref and LAP signals are digitised in A/D- converters 24 and 26 respectively before supplied to the microprocessor and supporting circuits 18. Data related to sensing and pacing, the measured impedance Z, and the activity or workload Act of the patient are supplied to the microprocessor and its supporting circuits 18 as well.

For communication with and transfer of data to external equipments the pacemaker is provided with telemetry means which are not shown in figure 3.

Figure 4 illustrates the determination of the quotient $\Delta P/\Delta V$. Since DHF evolves slowly, necessary measurements may be performed regularly, preferably 5 with long intervals, for a long time. A timer is therefore provided, block 28 in figure 4, and the process illustrated in the figure, is started by a check of status of the timer 28, block 30. The timer 28 may be set to e.g. hours, days or weeks.

As soon as the timer 28 has counted down to zero the measurements of LAP, Pref, the impedance Z corresponding to left ventricular volume, and the 10 activity or workload Act of the patient are started, block 32 in figure 4. It should be noted that the impedance is inversely correlated to the volume, but since the impedance variations are small and superimposed on a larger absolute impedance, impedance measurements are anyhow suitable for this purpose.

Measured data are stored in a ring buffer with n positions and large 15 enough to store data from pre-atrial loading of the left ventricle, block 34

As soon as a P-wave is detected, block 36, stored values of LAP, Pref, Z and Act are collected from a memory position n samples before the point of the P-wave detection. The number n depends on several circumstances, like the shape 20 of the P-wave, the filter in the P-wave amplifier, and the position of the right atrial electrode 12. In this way data are obtained from a point of time immediately before a noticeable change of the ventricular volume caused by the atrial contribution.

After this collection of parameter values average values of corresponding parameters, taken over several cardiac cycles, are created, block 38 in figure 4 and block 40 in figure 5. The average value creating process is illustrated in greater 25 detail in figure 5. The purpose of the average creation is to improve the quality of data.

The contribution from the atrium to the ventricular filling should not be part of the left ventricular volume measured in this procedure, as explained above. Thus data are extracted from a storage position n samples before the latest 30 sample, block 42 in figure 5, to avoid atrial contribution to the ventricular filling. Stored average values of LAP, Pref, Z and Act from previous heart beats are used, block 44, to produce a floating mean value by adding a fraction $1/k$ of the present value to the already existing "old" average value, according to the equation

$$\text{New_Average_Value} = \frac{\text{Value} + (k - 1) * \text{Old_Average_Value}}{k}$$

5 where "Value" means the present value.

This process is performed for each of the parameters LAP, Pref, Z and Act, block 46, and the new average values are stored, block 48 in figure 5. Block 46 contains corresponding equations for creating new average values of each of the parameters LAP, Pref, Z and Act, "avg" denoting "average".

10 It is then returned, block 50, to the main procedure illustrated in figure 3, where the next step consists in subtraction of Pref from LAP to obtain the corrected LAP value LAP_r as discussed above. Thus,

$$\text{LAP}_r = \text{LAP} - \text{Pref}$$

15

block 52 in figure 4. The corrected value LAP_r is then used to create the quotient ΔP/ΔV, block 54, store it in a memory, block 55, and compare the quotient ΔP/ΔV with a reference value for subsequent actions, block 56.

20 For this procedure the parameter values are preferably stored in an organized way, in this case in a table, see Table 1 below. This table is set up in a memory. The parameters are measured over time for different workload levels of the patient, and this is used to increase the number of available measurements to be used and to improve the results. When average values over several cardiac cycles of the parameters have been obtained LAP_r and Z are stored in a table 25 according to how actual activity or workload level fits the activity or workload intervals in the table. The following Table 1 shows an example of how to do this.

Activity interval	LAP _r	Z	Parameters set
Act ₁ to Act ₂	data	data	1
Act ₂ to Act ₃	no data	no data	0
...
Act _{n-1} to Act _n	data	data	1
N			Sum of above

The table will contain very few data at the beginning of the use of the apparatus according to the invention. With every entry into this process data will be filled in the table, either directly or via an averaging process if data already exist
5 in memory positions in question. An "1" in column "Parameters set" indicates that a true parameter has been obtained and stored in the table. An "0" indicates the opposite.

In Table 1 "N" denotes the number of activity or workload levels for which averaged parameter values have been stored and over which the quotient $\Delta P/\Delta V$
10 has to be determined.

For determining the quotient $\Delta P/\Delta V$, ΔP is first created for each indicated activity or workload interval in the table. All pressure data are referenced to the pressure for the patient in rest, Prest. The corresponding process is performed for ΔV . Since N activity or workload intervals are indicated as active in Table 1, N - 1
15 quotients $K_i = \Delta P_i/\Delta V_i$ are obtained. The average value of the K_i - values is returned, and $K_{avg} = (\sum K_i)/(N-1)$ is obtained. Thus the quotient K_{avg} is calculated.

The quotient K_{avg} is then compared to a predetermined reference value, preferably established and stored at the beginning of the use of the apparatus according to the invention (when the patient possibly were not yet suffering from
20 DHF). By storing subsequent quotients K_{avg} together with associated time marks the evolvement of the degree of DHF can be followed over time. A test can be set up to test whether the quotient has passed a certain limit for starting an automatic transmission of the DHF status of the patient to the physician. This message is transferred by means of the telemetry link of the apparatus according to the
25 invention.

CLAIMS

1. An implantable medical apparatus for detecting diastolic heart failure, DHF, comprising a DHF determining device for determining at least one blood pressure parameter for detecting a DHF state of the heart of a patient, **characterized in** that said DHF determining device comprises a pressure measuring means (2,26) for measuring the absolute value of left atrial pressure during the diastasis phase just before atrial contraction for a predetermined workload situation and a rest situation of the patient, and a comparison means (18) for comparing the difference, ΔP , between left atrial pressure measured in said workload and rest situations with a predetermined pressure difference reference value for DHF detection.
5
10. 2. An implantable medical apparatus for detecting diastolic heart failure, DHF, comprising a DHF determining device for determining at least one blood pressure parameter for detecting a DHF state of the heart of a patient, **characterized in** that said DHF determining device comprises a pressure measuring means for measuring the absolute value of the pressure in the pulmonary vein when the pulmonary valve is closed for a predetermined workload situation and a rest situation, and a comparison means (18) for comparing the difference, ΔP , between said pulmonary vein pressure measured in said workload and rest situations with a predetermined pressure difference reference value for DHF detection.
20
25. 3. The apparatus according to claims 1 or 2, **characterized in** that an activity sensor (22) is provided for determining the workload of the patient.
30. 4. The apparatus according to claim 1, **characterized in** that pressure compensating means (6,24) are provided for correcting the measured absolute value of left atrial pressure with the mean pressure in right atrium or in vena cava close to the right atrium.

5. The apparatus according to any of the preceding claims, **characterized in** that said DHF determining device comprises an impedance measuring means for measuring the impedance between a left ventricular electrode (8) and a second electrode (10) intended to be positioned in such a place that most of variations in the measured impedance are due to varying left ventricular volume, and in that ΔP values and corresponding values of the difference, ΔV , in the end-diastolic ventricular volume obtained from said impedances measured for said predetermined workload situation and for the patient in rest are supplied to a quotient forming means (54) to form the quotient $\Delta P/\Delta V$, and in that said comparison means (56) is adapted to compare said quotient $\Delta P/\Delta V$ with a predetermined quotient reference value for DHF detection.

10

15. 6. The apparatus according to any of the preceding claims, **characterized in** that a first averaging means is provided to form an average value of said absolute pressures measured during a plurality of cardiac cycles with said predetermined workload situation and an average value of said absolute pressures measured during a plurality of cardiac cycles with the patient in rest.

20

25. 7. The apparatus according to claim 6, **characterized in** that a second averaging means is provided to form an average value of said impedance measured during a plurality of cardiac cycles with said predetermined workload situation and an average value of said impedance measured during a plurality of cardiac cycles with the patient in rest.

30. 8. The apparatus according to any of the preceding claims, **characterized in** that a wireless communication means is connected to said comparison means for automatically sending the results of the comparison of said difference between measured pressures with said pressure difference reference values to external receiver means.

9. The apparatus according to claim 8, **characterized in** that said wireless communication means is connected to said comparison means for automatically sending the results of the comparison of said difference between said quotient $\Delta P/\Delta V$ in workload and rest situations with said quotient reference value to said external receiver means.

10. The apparatus according to any of the claims 1 - 7, **characterized in** that a storing means is provided for storing the results of the comparison of said difference between measured pressures with said pressure difference reference values.

11. The apparatus according to claim 10, **characterized in** that said storing means is provided for storing the results of the comparison of said difference between said quotient $\Delta P/\Delta V$ in workload and rest situations with said quotient reference value.

12. A pacemaker, **characterized in** that it comprises an apparatus according to any one of the preceding claims and control means (14) for optimising pacing therapy depending on the result of said comparisons with said predetermined reference values.

13. The pacemaker according to claim 12, **characterized in** that it comprises a rate responsive sensor (22) for determining the workload situation of the patient.

14. The pacemaker according to claim 12 comprising a pressure sensor, **characterized in** that said pressure sensor (22) is arranged as said activity sensor.

15. The pacemaker according to any of the claims 12 – 14, **characterized in** that it comprises pressure compensating means including a pressure sensor located on the pacemaker capsule or on a satellite, intended to be implanted in body tissue close to the pacemaker.

16. A method of detecting diastolic heart failure, DHF, comprising the step of determining at least one blood pressure parameter for detecting a DHF state of the heart of a patient, **characterized in** that absolute value of left atrial pressure is measured (32) during the diastasis phase just before atrial contraction for a predetermined workload situation and a rest situation of the patient, and the difference, ΔP , between left atrial pressure measured in said workload and rest situations is compared (56) with a predetermined pressure difference reference value for DHF detection.

17. A method of detecting diastolic heart failure, DHF, comprising the step of determining at least one blood pressure parameter for detecting a DHF state of the heart of a patient, **characterized in** that absolute value of the pressure in the pulmonary vein is measured when the pulmonary valve is closed for a predetermined workload situation and a rest situation of the patient, and the difference, ΔP , between said pulmonary vein pressure measured in said workload and rest situations is compared with a predetermined pressure difference reference value for DHF detection.

18. The method according to claims 16 or 17, **characterized in** that the impedance is measured between a left ventricular electrode (8) and a second electrode (10) intended to be positioned in such a place that most of variations in the measured impedance are due to varying left ventricular volume for said predetermined workload situation and for the patient in rest, whereupon the end-diastolic volume is determined from a predetermined relation between said measured impedance and the left ventricular volume, and in that a quotient between said difference between measured absolute pressure values, ΔP , and the corresponding difference between end-diastolic volumes, ΔV , for said workload and said rest situations, is formed (54) and compared (56) with a predetermined quotient reference value for DHF detection.

19. The method according to claim 18, **characterized in** that in a calibration procedure instantaneous values of the left ventricular volume are measured by ultrasound technique and said relation is established between these measured ventricular volume values and simultaneously measured impedance values.
5
20. The method according to any of the claims 17 - 20,**characterized in** that an average value of said absolute pressures is measured (38,40,42,44,46) during a plurality of cardiac cycles with said predetermined workload situation and an average value of said absolute pressures is measured during a plurality of cardiac cycles with the patient in rest.
10
21. The method according to claim 20, **characterized in** that an average value of said impedance is measured (38,40,42,44,46) during a plurality of cardiac cycles with said predetermined workload situation and an average value of said impedance is measured during a plurality of cardiac cycles with the patient in rest.
15
22. The method according to any of the claims 16– 21, **characterized in** that the results of the comparison of said pressure differences with said pressure difference reference values are automatically sent to external receiver means.
20
23. The method according to claim 22, **characterized in** that the results of the comparison (56) of said difference between said quotient $\Delta P/\Delta V$ in workload and rest situations with said quotient reference value are automatically sent to external receiver means.
25
24. The method according to claim 16, **characterized in** that the measured absolute value of left atrial pressure is corrected with the mean pressure in right atrium or in vena cava close to the right atrium.
30

25. The method according to any of the claims 16 – 24, **characterized in** that
said pressures are measured for different workloads of the patient and
5 for the patient in rest at an early time, when the patient is not suffering
from DHF, for determining said pressure difference reference values.

26. The method according to claim 25, **characterized in** that said impedance
is measured for different workloads of the patient and for the patient in
10 rest at an early time, when the patient is not suffering from DHF, for
determining said quotient reference value.

ABSTRACT

An implantable medical apparatus for detecting diastolic heart failure, DHF, comprises a DHF determining device for determining at least one blood pressure parameter for detecting a DHF state of the heart (4) of a patient. The DHF determining device comprises a pressure measuring means (2,26) for measuring the absolute value of left atrial pressure during the diastasis phase just before atrial contraction, or the absolute value of the pressure in the pulmonary vein when the pulmonary valve is closed, for a predetermined workload situation and a rest situation of the patient. A comparison means (18) compares the difference between left atrial pressure or pulmonary vein pressure, in said workload and rest situations, with a predetermined pressure difference reference value. A pacemaker comprises such an apparatus and control means (14) for optimising pacing therapy depending on the result of the comparisons with the reference values. A corresponding method of detecting diastolic heart failure, DHF, comprises the step of determining at least one blood pressure parameter for detecting a DHF state of the heart of a patient. The absolute value of left atrial pressure is measured during the diastasis phase just before atrial contraction, or the absolute value of the pressure in the pulmonary vein when the pulmonary valve is closed, for a predetermined workload situation and a rest situation of the patient. The difference between left atrial pressure or pulmonary vein pressure in said workload and rest situations is compared with a predetermined pressure difference reference value.

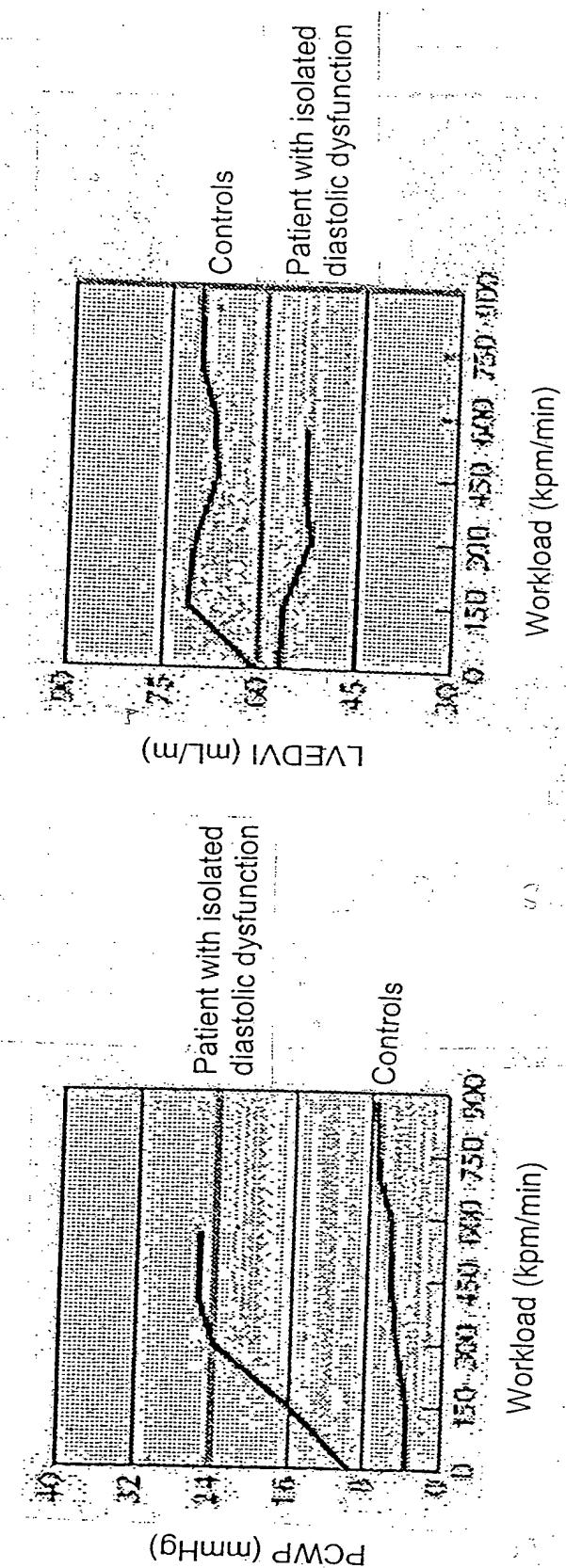


Fig. 1

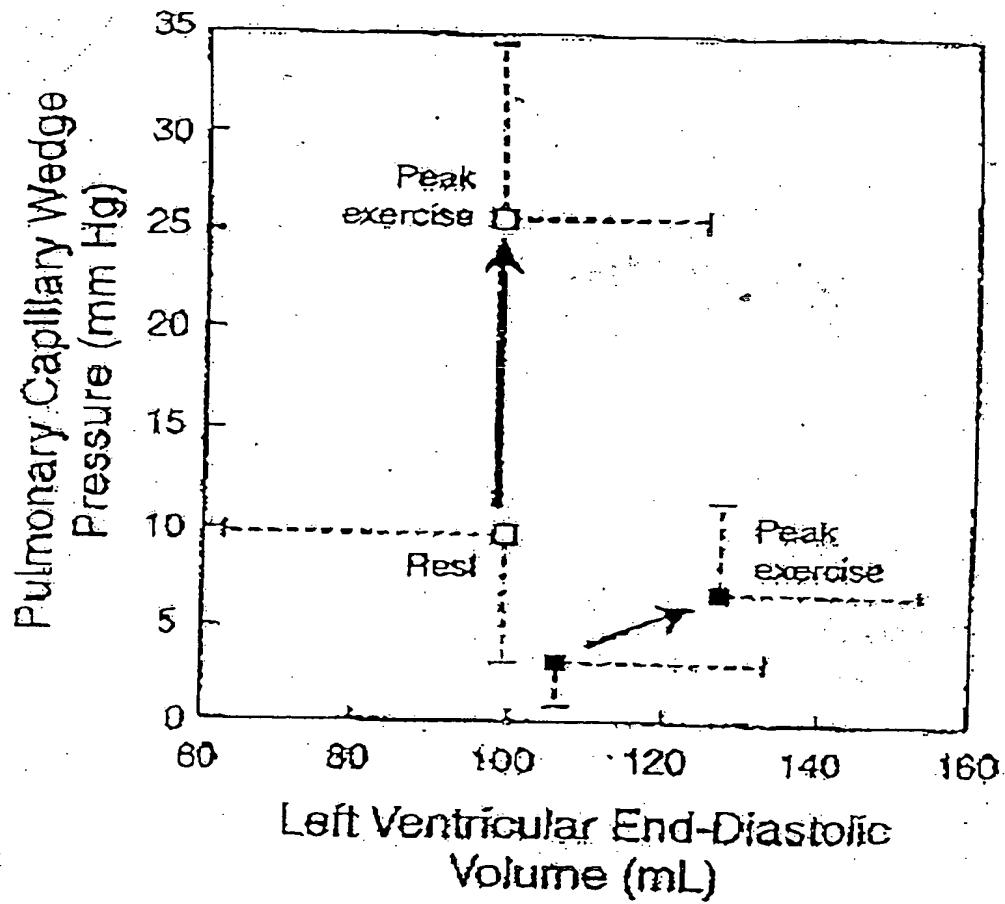


Fig. 2

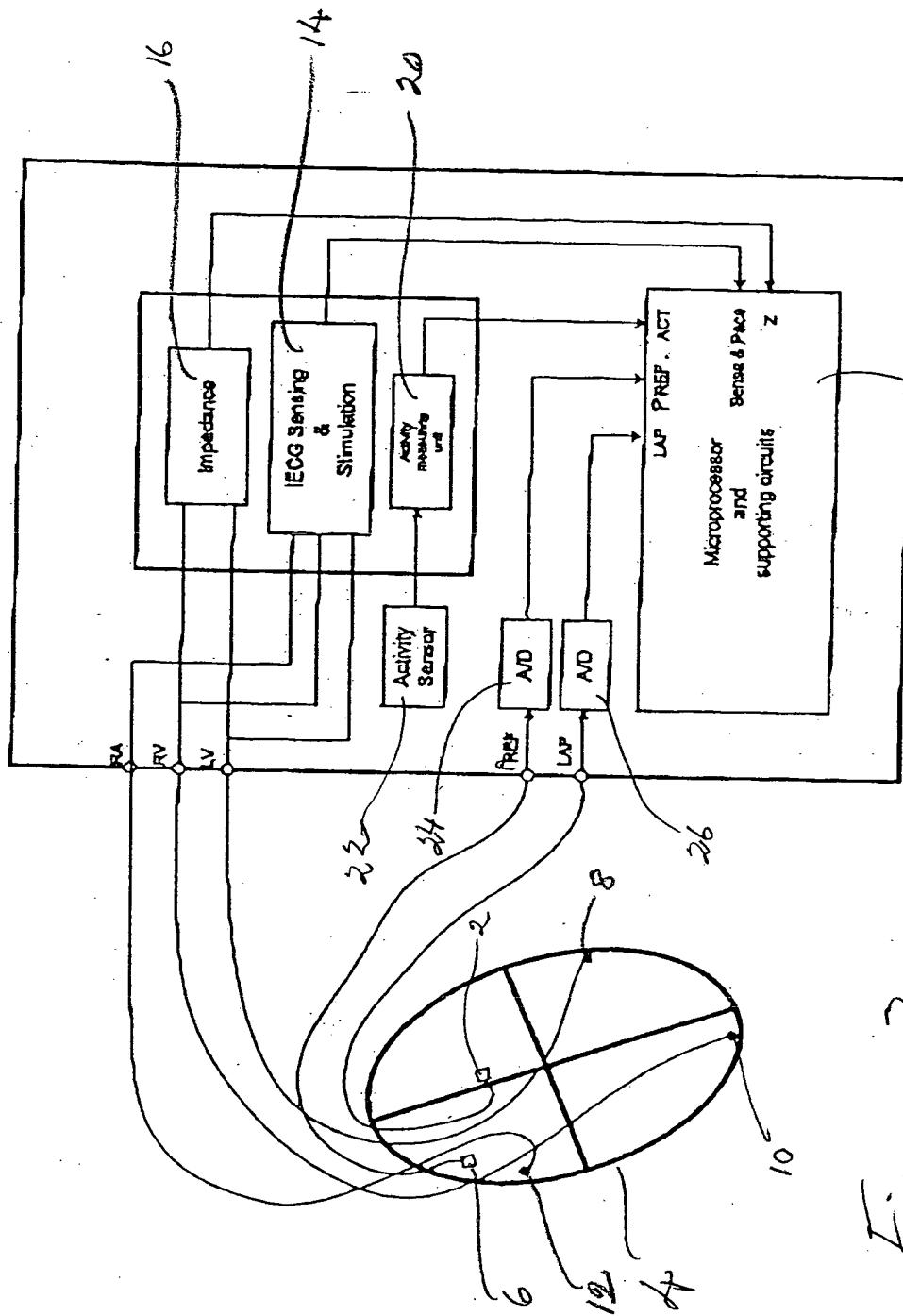


Figure 3

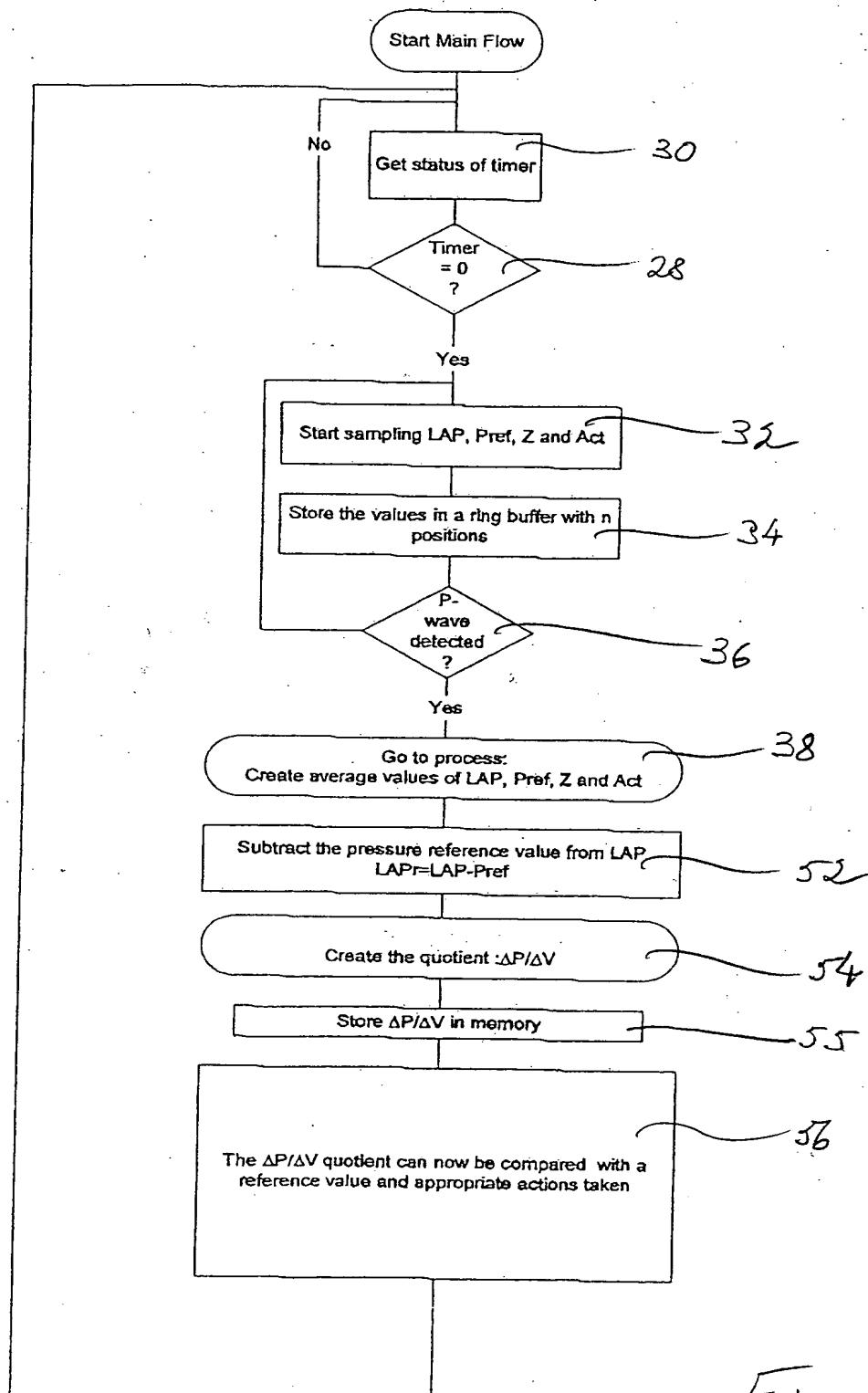


Fig. 4

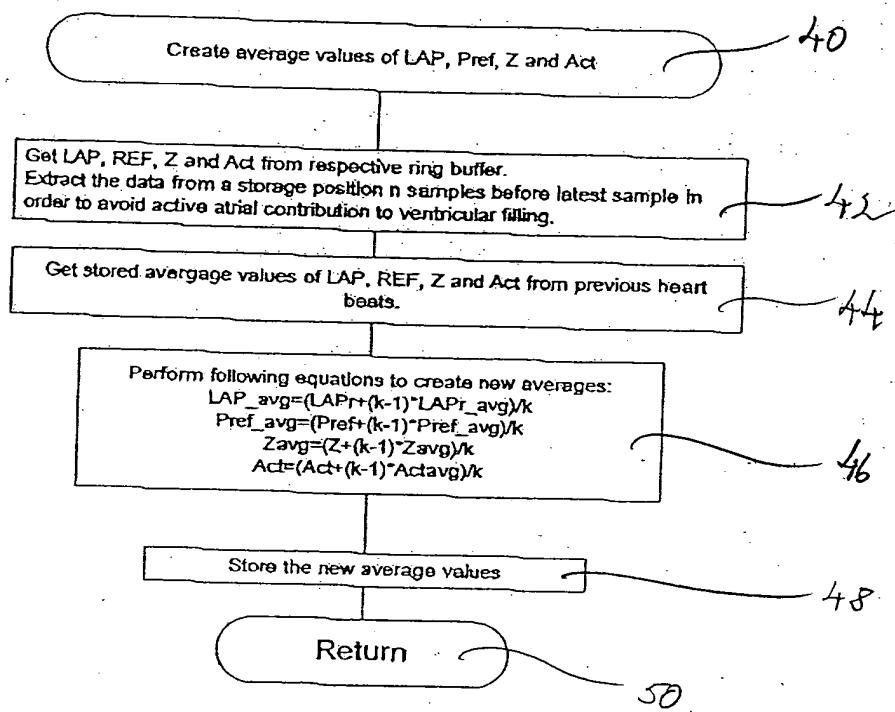


Fig. 5